

Two Dimensional Linkage Study Methods and Related Inventions

2 The present patent application is a continuation-in-part of U.S. Patent Application 09/947,768 (filed 5
3 SEPT 2001). And 09/947,768 claims priority from US Provisional 60/230570 (filed 9/5/2000). Patent
4 Application 09/947,768 is a continuation-in-part of U.S. Patent Application 09/623,068 (filed 26 AUG
5 2000). The present patent application is also, a continuation-in part of Patent Application 09/623,068
is a 371 of and
6 (filed 26 AUG 2000). Application 09/623,068 claims priority from PCT/US99/04376 filed (2/26/99).

7 PCT/US99/04376 claims priority from US Provisional applications: 60/076182 filed 27 Feb 1998,
8 60/086947 filed 27 May 1998, 60/076102 filed 26 Feb 1998 and 60107673 filed 7 Nov 1998. Each of the
9 following patent applications are incorporated herein by reference in their individual entireties: U.S.
10 Provisional Patent Application 60/230570, PCT/US99/04376, U.S. Patent Application 09/623,068, and
11 U.S. Patent Application 09/947,768.

12 The reader's attention is directed to the following documents or papers each of which is open to the
13 public and each of which is incorporated by reference herein in their entirety: (1) McGinnis, Ewens &
14 Spielman, Genetic Epidemiology 1995 ; 12(6) : 637-40. (2) RE McGinnis Annals of Human Genetics vol
15 62, pp. 159-179

Technical Field

16 Versions of the present invention are in the field of molecular biology, some versions are specifically in
17 the area of finding the chromosomal location of genes that cause genetic characteristics such as
18 human disease.

Background

Introduction

21 Conventional linkage study techniques have limited power to localize trait causing genes (trait causing
22 polymorphisms) of modest effect, such as many human disease polymorphisms. The two-dimensional
23 linkage study techniques of this application are powerful new techniques for localizing genes
24 (polymorphisms) especially of modest effect.

Chromosomes, heredity, genes, markers and alleles

25 Chromosomes are large molecules that carry the information for the inheritance of physical (genetic)
26 characteristics or traits. In human beings for example, parents pass a copy of half of their chromosomes
27 to their offspring during reproduction. By doing this, each parent passes some of his or her physical
28 characteristics to his or her offspring. Any chromosome of a living creature is made of a large string-like
29 molecule of DNA. Chromosomes are essentially very long strings of DNA. Genes are small pieces of a
30 chromosome that cause or determine inherited genetic characteristics. (In this application, the term
31 gene means a polymorphism that determines a genetic characteristic; the term does not mean an entire
32 gene structure with a promoter region, introns, etc..) Markers are any segment of DNA on a
33 chromosome which can be identified and whose chromosomal location is known (at least to some
34 extent). Markers are like milestones along the very long string-like molecule of DNA which makes up a
35 chromosome. Both a gene and a marker can come in different forms on different chromosomes. These